Prisca	5.0.2.37
Date of report:	**13-03-2019**

Patient data

Name	MRS. PREETI JOSHI	Patient ID	1903220768/AMB
Birthday	01-01-1992	Sample ID	1903220768/AMB
Age at sample date	27.2	Sample Date	13-03-2019
Gestational age	12 + 3		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregancies	no
Weight	56	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data / Ultrasound data

Parameter	Value	Corr. MoM	Ultrasound data	
			Gestational age	12 + 0
PAPP-A	2.47 mIU/ml	0.68	Method	CRL Robinson
fb-hCG	48.6 ng/ml	1.32	Scan date	10-03-2019

Risks at sampling date

Age risk	1:845	Crown rump length in mm	56.8
Biochemical T21 risk	1:1159	Nuchal translucency MoM	0.60
Combined trisomy 21 risk	1:6625	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	.
		Qualifications in measuring NT	MD

Risk

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 6625 women with the same data, there is one woman with a trisomy 21 pregnancy and 6624 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.

Please note that risk calculations are statistical approaches and have no diagnostic value!

The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).

The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Sign of Physician

below cut off	Below Cut Off, but above Age Risk	above cut off